PTO/SB/08s (08-03)
Approved for use through 97/31/2006, ONE 0651-0231
U.S. Patent and Trademark Office, U.S. DEPARTMENT OF COMMERCE to a collection of information unless it contains a valid OMB control number. Under the Paperwork Reduction Act of 1995, no persons are required to

U.S.PATENTS

INFORMATION DISCLOSURE
STATEMENT BY APPLICANT
(Not for submission under 37 CFR 1.99)

Application Number		10541042
Filing Date		2005-06-24
First Named Inventor Shiny		a Adachi
Art Unit		3661
Examiner Name BEAU		ILIEU, YONEL
Attorney Docket Number		38404

Examiner Initial*	Cite No	Patent Number	Kind Code ¹	Issue E)ate	Name of Patentee or Applicant of cited Document		Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear		
	1									
If you wis	h to a	i dd additional U.S. Patei	nt citatio	n inform	ation pl	l lease click the	Add button.		Add	
			U.S.P	ATENT	APPLI	CATION PUB	LICATIONS		Remove	
Examiner Initial*	Cite No	Publication Number	Kind Code ¹	Publica Date	tion	of cited Decument		Releva	s,Columns,Lines where ant Passages or Releases Appear	e vant
	1									
If you wis	h to a	dd additional U.S. Publi	shed Ap	plication	citatio	n information p	olease click the Ad	d buttor		
				FOREK	SN PA1	ENT DOCUM	IENTS		Remove	
Examiner Initial*	Cite No	Foreign Document Number ³	Country Code ²		Kind Code ⁴	Publication Date	Name of Patente Applicant of cited Document	e or	Pages,Columns,Lines where Relevant Passages or Relevant Figures Appear	TE
	1	10-283591	JP			1998-10-23				×
	2	2000-180193	JP			2000-06-30				×
	3	2002-533854	JP			2002-10-08				×
			1			1				_

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 CFR 1.99)

Application Number		10541042
iling Date		2005-06-24
irst Named Inventor	Shiny	a Adachi
Art Unit		3661
Examiner Name BEAU		ILIEU, YONEL
Marnov Docket Number		28404

If you wish	to a	id add	ditional Foreign Patent Document citation information please click the Add bu	tton Add	
			NON-PATENT LITERATURE DOCUMENTS	Remove	
Examiner Initials*	Examiner Cite Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc), date, pages(s), volume-issue number(s), publisher, city and/or country where published.				
	1				
If you wish	n to a	id add	fitional non-patent literature document citation information please click the A	d button Add	
			EXAMINER SIGNATURE		
Examiner	Signa	ture	Date Considered	1	
			reference considered, whether or not citation is in conformance with MPEP i rmance and not considered. Include copy of this form with next communicat		

1 See Kind Code of USFTO Planto Documents at year, <u>USFTO CODY</u> or MEPD 910.4. ² Enter office that issued the cocument, by the two-letter code (WPD Standard ST3.). ² The Japanese parted coverents, the relication to the year of the region of the Emperor must proceed the serial number of the papert document of the Standard ST3.0 ² Serial ST3.0 ² Serial Standard ST3.0 ² Serial ST3.0 ² Serial Standard ST3.0 ² Serial ST3.0 ² S

INFORMATION DISCLOSURE STATEMENT BY APPLICANT (Not for submission under 37 GFR 1.99)

Application Number		10541042
Filing Date		2005-06-24
First Named Inventor Shiny		a Adachi
Art Unit		3661
Examiner Name BEAU		LIEU, YONEL
Attorney Docket Number		38404

CERTIFICATION STATEMENT

Please see 37 CFR 1.97 and 1.98 to make the appropriate selection(s):

	That each item of information contained in the information disclosure statement was first cited in any communication
X	from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of the
	information displacate statement. Con 27 CED 1 07(a)/1)

OR

That no item of information contained in the information disclosure statement was cited in a communication from a
foreign patent office in a counterpart foreign application, and, to the knowledge of the person signing the certification
after making reasonable inquiry, no item of information contained in the information disclosure statement was known to
any individual designated in 37 CFR 1.56(c) more than three months prior to the filing of the information disclosure

- See attached certification statement.
- Fee set forth in 37 CFR 1.17 (p) has been submitted herewith.
-

SIGNATURE

A signature of the applicant or representative is required in accordance with CFR 1.33, 10.18. Please see CFR 1.4(d) for the form of the signature.

Signature	/jeffrey j sopko/	Date (YYYY-MM-DD)	2007-01-03
Name/Print	Jeffrey J. Sonko	Registration Number	27676

This collection of information is required by 3T CFR 1.97 and 1.98. The information is required to obtain or retain a benefit by the public which is for life railed by the USPTO to process) an application. Confidentiality is governed by \$5 U.S. C. 12.0 and 3T CFR 1.14. This collection is estimated to take 1 hour to complete, including gathering, preparing and submitting the completed application from the USPTO. Time will vary depending upon the individual case. Any comments on the amount of time you require to complete his form and/or suggestions for reducing this burden, should be sent to the Chief Information Officer, U.S. Patent and Tradenar's Office, U.S. operatment of Commence, P. 0. Box 1450, Alexandria, V.S. 2313-1450. DIN OTS CRID FEES OR COMPLETED FORMS TO THIS ADDRESS. SEND TO: Commissioner for Patents, P.O. Box 1450, Alexandria, V.S. 2313-1450.

Privacy Act Statement

The Privacy Act of 1974 (P. L. 93-579) requires that you be given certain information in connection with your submission of the stackhold from related to a patient application or patient. Accordingly, pursuant to the requirements of the Act, places be advised that (1) the general authority for the collection of this information is 35 U.S.C. 2(b)(2); (2) familishing of the information solicided is couldrain; and (3) the primoral pursuance for which the information is used by the U.S. Patient and Trademan Coffice is to process and/or examine your submission related to a patient agricultant or patient. If you do not furnish the requested process and/or examine your submission related to a patient agricultant or patient. If you do not furnish the requested results of the patient of the patient and the patient of the patient

The information provided by you in this form will be subject to the following routine uses:

- The information on this form will be treated confidentially to the extent allowed under the Freedom of Information Act (5 U.S.C. 552) and the Privacy Act (5 U.S.C. 552a). Records from this system of records may be disclosed to the Department of Justice to determine whether the Freedom of Information Act requires disclosure of these record s.
 - A record from this system of records may be disclosed, as a routine use, in the course of presenting evidence to a court, magistrate, or administrative tribunal, including disclosures to opposing counsel in the course of settlement negotiation.
 - A record in this system of records may be disclosed, as a routine use, to a Member of Congress submitting a request involving an individual, to whom the record pertains, when the individual has requested assistance from the Member with respect to the subject matter of the record.
 - A record in this system of records may be disclosed, as a routine use, to a contractor of the Agency having need for the information in order to perform a contract. Recipients of information shall be required to comply with the requirements of the Privacy Act of 1974, as amended, pursuant to 5 U.S.C. 552(m).
 - A record related to an International Application filed under the Patent Cooperation Treaty in this system of records
 may be disclosed, as a routine use, to the International Bureau of the World Intellectual Property Organization, pursuant
 to the Patent Cooperation Treaty.
 - A record in this system of records may be disclosed, as a routine use, to another federal agency for purposes of National Security review (35 U.S.C. 181) and for review pursuant to the Atomic Energy Act (42 U.S.C. 218(c)).
 - 7. A record from this system of records may be disclosed, as a routine use, to the Administrator, General Services, or hisher designed, uturing an insection of records conducted by GSA as part of that agency's responsibility to recommend improvements in records management practices and programs, under authority of 4d U.S.C. 2904 and 2905. Such disclosure shall be made in accordance with the GSA requisions governing inseption of records for this purpose, and any other relevant (i.e., GSA or Commerce) directive. Such disclosure shall not be used to make determinations about individuals.
- A record from this system of records may be disclosed, as a routine use, to the public after either publication of the
 application pursuant to 35 U.S.C. 12(2) or issuance of a patent pursuant to 35 U.S.C. 151. Further, a record may be
 disclosed, subject to the limitations of 37 CFR 1.14, as a routine use, to the public if the record was filled in application
 which became abandoned or in which the proceedings were terminated and which application is referenced by either a
 published application, an application open to public inspections or as issued patent.
 - A record from this system of records may be disclosed, as a routine use, to a Federal, State, or local law enforcement agency, if the USPTO becomes aware of a violation or potential violation of law or regulation.